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STRUCTURE FILE UPDATES: 9 JAN 2004 HIGHEST RN 635758-32-6 DICTIONARY FILE UPDATES: 9 JAN 2004 HIGHEST RN 635758-32-6

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Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

=> s gamma aminobutyramide

122714 GAMMA

25 AMINOBUTYRAMIDE

L1 3 GAMMA AMINOBUTYRAMIDE

(GAMMA (W) AMINOBUTYRAMIDE)

=> d 11 1-3

L1 ANSWER 1 OF 3 REGISTRY COPYRIGHT 2004 ACS on STN

RN 35821-20-6 REGISTRY

CN Carbamic acid, (4-amino-4-oxobutyl)-, phenylmethyl ester (9CI) (CA INDEX NAME)

OTHER NAMES:

CN N-(Benzoxycarbonyl)-γ-aminobutyramide

CN N-Benzyloxycarbonyl-γ-aminobutyric acid amide

FS 3D CONCORD

MF C12 H16 N2 O3

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT, IFICDB, IFIPAT, IFIUDB, TOXCENTER, USPATFULL

(*File contains numerically searchable property data)

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

9 REFERENCES IN FILE CA (1907 TO DATE)

9 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L1 ANSWER 2 OF 3 REGISTRY COPYRIGHT 2004 ACS on STN

RN 25271-48-1 REGISTRY

CN Hydrocinnamamide, β -(aminomethyl)-, dihydrochloride, DL-(8CI) (CA INDEX NAME)

OTHER NAMES:

CN DL-β-Phenyl-γ-aminobutyramide dihydrochloride

MF C10 H14 N2 O . 2 Cl H

LC STN Files: CA, CAPLUS, TOXCENTER

CRN (102571-00-6)

2 HC1

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L1 ANSWER 3 OF 3 REGISTRY COPYRIGHT 2004 ACS on STN

RN 3459-33-4 REGISTRY

CN 2H-Isoindole-2-butanamide, 1,3-dihydro-1,3-dioxo- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

N 2-Isoindolinebutyramide, 1,3-dioxo- (7CI, 8CI)

OTHER NAMES:

4-Phthalimidobutyramide

N-Phthalyl-γ-aminobutyramide

FS 3D CONCORD

C12 H12 N2 O3 MF

BEILSTEIN*, CA, CAOLD, CAPLUS, CHEMCATS, RTECS*, TOXCENTER, LC STN Files: USPAT2, USPATFULL

(*File contains numerically searchable property data)

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 12 REFERENCES IN FILE CA (1907 TO DATE)
- 1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
- 12 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- 2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> file medicine

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PATENT INFORMATION:

US 2003100933

APPLICATION INFO .: RELATED APPLN. INFO.:

20020924 (10) A1US 2002-254024 Continuation-in-part of Ser. No. US 2001-843334, filed on 26 Apr 2001, PENDING Continuation-in-part of Ser. No. US 2002-205474, filed on 24 Jul 2002, PENDING Continuation-in-part of Ser. No. WO 2002-IL68, filed on 23 Jan 2002, UNKNOWN Continuation-in-part of Ser. No. US 2001-944913, filed on 31 Aug 2001, PENDING

NUMBER DATE ______

PRIORITY INFORMATION:

US 2002-383157P 20020523 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE: DARBY & DARBY P.C., P. O. BOX 5257, NEW YORK, NY,

10150-5257

NUMBER OF CLAIMS:

250

EXEMPLARY CLAIM: NUMBER OF DRAWINGS:

7 Drawing Page(s)

LINE COUNT:

SUMM

[0007] Cerebral palsy (CP) is an upper motor neuron (UMN) disorder typically caused by brain injury at or around the time of birth, or in the first year of an infant's life. The most common type of CP is spastic CP, which causes spasticity, mainly of the flexor muscles of the arms and legs. In CP, spasticity is caused by.

SUMM

. . . and clonazepam), dantrolene, and, more recently, tizanidine (Zanaflex). Additionally, recent studies have shown that injecting Botox (the Botulinum toxin) into spastic muscles can bring relief by causing the muscles to relax.

SUMM

. . . PCT Patent Publication WO 01/10432 to Meythaler et al., which is incorporated herein by reference, describes a method for treating spastic disorders, convulsive disorders, pain and epilepsy by administering a therapeutically-effective amount of the compound gamma-aminobutyramide and analogs thereof.

SUMM

. . . the constants are pre-set for a given condition. The physiological parameters are preferably indicative of the onset or strength of spastic muscle contraction and/or limb movement. The control unit preferably comprises a sensor unit for measuring the physiological parameters. For some.

DETD

. . . the constants are pre-set for a given condition. The physiological parameters are preferably indicative of the onset or strength of spastic muscle contraction and/or limb movement. For some applications, control unit 20 drives the electrode device to apply a current in.

ANSWER 2 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN L7

Citing Full References Text ACCESSION NUMBER:

2001:114975 CAPLUS

DOCUMENT NUMBER:

134:141767

TITLE:

Use of GABA agonists for treatment of spastic disorders, convulsions, epilepsy, and pain

INVENTOR(S):

Meythaler, Jay M.; Peduzzi, Jean UAB Research Foundation, USA

PATENT ASSIGNEE(S):

PCT Int. Appl., 28 pp.

SOURCE: CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

APPLICATION NO. DATE KIND DATE PATENT NO. _ - - - - - - -WO 2000-US21886 20000810 WO 2001010432 A1 20010215

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,

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SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
             YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
             CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                           EP 2000-955436
                                                             20000810
                           20020508
     EP 1202720
                       A1
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL
                                            JP 2001-514950
                                                             20000810
     JP 2003506407
                            20030218
                      T2
                                                             20000810
                            20031031
                                            NZ 2000-517407
     NZ 517407
                       Α
PRIORITY APPLN. INFO.:
                                         US 1999-148159P P 19990810
                                         WO 2000-US21886 W 20000810
                               THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
     Use of GABA agonists for treatment of spastic disorders, convulsions,
TI
     epilepsy, and pain
     A method is disclosed for treating a patient/subject having a spastic
AB
     disorder, a convulsive disorder, pain, or epilepsy which includes
     administering a therapeutically effective amt. of \gamma-
     aminobutyramide, analogs, substituted forms, derivs., pharmaceutically
     acceptable salts, esters, amides, and prodrugs thereof, or compds. which
     yield \gamma-aminobutyramide as an intermediate, metabolite, or
     byproduct.
     GABA agonist spastic disorder convulsion epilepsy pain;
ST
     aminobutyramide spastic disorder convulsion epilepsy pain
     Medical goods
IT
        (catheters, spinal; γ-aminobutyramide and
        related compds. for treatment of spastic disorders,
        convulsions, epilepsy, and pain)
IT
     Nervous system
        (dystonia; y-aminobutyramide and related
        compds. for treatment of spastic disorders, convulsions,
        epilepsy, and pain)
IT
     Muscle, disease
        (hypertonia, spastic; γ-
        aminobutyramide and related compds. for treatment of
        spastic disorders, convulsions, epilepsy, and pain)
IT
     Drug delivery systems
        (implants, pumps; γ-aminobutyramide and
        related compds. for treatment of spastic disorders,
        convulsions, epilepsy, and pain)
     Drug delivery systems
IT
        (intrathecal; γ-aminobutyramide and related
        compds. for treatment of spastic disorders, convulsions,
        epilepsy, and pain)
IT
     Drug delivery systems
        (intraventricular; γ-aminobutyramide and
        related compds. for treatment of spastic disorders,
        convulsions, epilepsy, and pain)
IT
     Drug delivery systems
        (prodrugs; y-aminobutyramide and related
        compds. for treatment of spastic disorders, convulsions,
        epilepsy, and pain)
IT
     Nervous system
        (spasticity; y-aminobutyramide and related
        compds. for treatment of spastic disorders, convulsions,
        epilepsy, and pain)
     Brain, disease
IT
        (trauma; y-aminobutyramide and related
        compds. for treatment of spastic disorders, convulsions,
        epilepsy, and pain)
IT
     Analgesics
     Anticonvulsants
        (y-aminobutyramide and related compds. for
        treatment of spastic disorders, convulsions, epilepsy, and
```

~ pain)

IT $\frac{1134-47-0}{3251-08-9}$, γ -Aminobutyric acid amide $\frac{3251-08-9}{1120}$, γ -Aminobutyric acid amide, derivs. $\frac{62666-20-0}{1120}$,

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

 $(\gamma\text{-aminobutyramide}$ and related compds. for treatment of spastic disorders, convulsions, epilepsy, and pain)

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Full Citing Text References

on STN

ACCESSION NUMBER: 80219761 EMBASE

DOCUMENT NUMBER: 1980219761

TITLE: New anticonvulsants: Schiff bases of γ-aminobutyric

acid and y-aminobutyramide.

AUTHOR: Kaplan J.P.; Raizon B.M.; Desarmenien M.; et al.

CORPORATE SOURCE: Dept. Chem., CNS Unit, Lab. Etudes Rech. SYNTHELABO, 92220

Bagneux, France

SOURCE: Journal of Medicinal Chemistry, (1980) 23/6 (702-704).

CODEN: JMCMAR United States

COUNTRY: United : DOCUMENT TYPE: Journal

FILE SEGMENT: 050 Epilepsy

LANGUAGE: English

TI New anticonvulsants: Schiff bases of γ -aminobutyric acid and

γ-aminobutyramide.

AB Schiff bases of γ-aminobutyric acid (γAbu) and

γ-aminobutyramide (γAbuNH2) were prepared and tested for

anticonvulsant and γAbu mimetic activity. 4-[[(4-Chlorophenyl)(5-

fluoro-2-hydroxyphenyl) methylene] amino] butanoic acid monosodium salt (4) and 4-[[(4-chlorophenyl) (5-fluoro-2-hydroxyphenyl) methylene] amino] butanami

de (5) blocked.

CT Medical Descriptors:

*4 aminobutyric acid h 3

brain

cerebellum

convulsion

drug distribution

structure activity relation

central nervous system

in vitro study

animal experiment

mouse rat

normal human

*progabide acid

*bicuculline

*progabide

*muscimol

*phenytoin

*valproic acid

calcium 45

L7 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2004 ACS on STN

Full Citing
Text References

ACCESSION NUMBER: 1975:557909 CAPLUS

DOCUMENT NUMBER: 83:157909

TITLE: Elevation of brain γ -aminobutyric acid in the

mouse by L-2,4-diaminobutyric acid

AUTHOR(S): Shuter, Eli R.; Robins, Eli

CORPORATE SOURCE:

St. Louis, MO, USA

SCURCE:

Transactions of the American Neurological Association

(1974), 99, 49-51

CODEN: TANAA4; ISSN: 0065-9479

DOCUMENT TYPE:

Journal

English LANGUAGE: In mice injected with L-2,4-diaminobutyric acid (I) [1758-80-1] (50 mg) AΒ and sacrificed at 1, 10, and 20 hr or allowed to die, there was a marked increase in brain GABA [56-12-2]; after 1 hr, 8.3%; after 10 hr, 58.4%;

after 20 hr, 94.8%, and at time of death 153%. Et γ -aminobutyrate-HCl (II) [6937-16-2] and γ -aminobutyryl choline chloride-HCl (III)

[2922-18-1] produced convulsions, and \(\gamma \)-aminobutyramide [5959-32-0] produced sedation, all without altering brain GABA content.

II and III may block GABA receptors at inhibitory synapses, whereas I may prevent the uptake of GABA at presynaptic storage sites following synaptic release.

2922-18-1 6937-16-2 IT

RL: BIOL (Biological study)

(GABA of brain in response to, convulsion in relation to)